Central Line-Associated Bloodstream Infection (CLABSI)
NHSN CLABSI protocol http://www.cdc.gov/nhsn/PDFs/pscManual/4PSC_CLABScurrent.pdf

		Date of admission		
Date of central line insertion	aI	Date of central line removal	Date of review	Date of event
criteria may be present d	uring the first 2 calend	sent together ≥ 3 calendar days of adar days of admission as long as they cur within a timeframe that does not	are also present on or after c	alendar day 3 of admission
with day of device places		neter (UC) in place for > 2 calendar of the patient is admitted or transferred 3 of line		
AND				
☐ A CL or UC is in place of Line in place on d	•	day before ine in place day before event		
Patient must meet at least o	ne of the following cri	teria for laboratory-confirmed blo	odstream infection (LCBI):	
Criterion 1 (for any patier Patient has a recognized AND Organism cultured from	pathogen cultured from	m one or more blood cultures (see no	otes 1 and 2)	
organism cultured from	blood is not related to	an infection at another site		
Criterion 2 (for any patient) □ Patient has at least one of office (> 38°) □ chills □ hypotension AND	f the following signs o	r symptoms:		
□ Positive laboratory result	s are <u>not</u> related to an	infection at another site		
AND				
spp., coagulase-negative cultured from two or mo	e staphylococci (includore blood cultures drav	oids (<i>Corynebacterium</i> spp. not <i>C. di</i> ding <i>S. epidermidis</i>), viridans group wn on separate occasions. Criterion of mmon commensals at http://www.co	streptococci, Aerococcus spp elements must occur within a	., and <i>Micrococcus</i> spp.] is timeframe that does not exceed a





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Patient name	MR #	Date of admission		
Date of central line inse	ertion	Date of central line removal		Date of event
criteria may be prese	ent during the first 2	rst present together ≥ 3 calendar days of a calendar days of admission as long as the cria occur within a timeframe that does no	ey are also present on or after	calendar day 3 of admission
with day of device p □ Day 1 of line		cal catheter (UC) in place for > 2 calendary 1(or, if the patient is admitted or transfer Day 3 of line		<u> </u>
AND □ A CL or UC is in pla	ace on the day of ev	ent or day before		
	•	☐ Line in place day before event		
 fever (> hypothe apnea bradyca AND Positive laboratory r 	ge has at least <u>one</u> of 38° C core) rmia (< 36° C core) rdia	of the following signs or symptoms: d to an infection at another site		
coagulase-negative s two or more blood c	staphylococci (inclu ultures drawn on se	(Corynebacterium spp. not C. diphtherial ding S. epidermidis), viridans group strep parate occasions. Criterion elements must non commensals at http://www.cdc.gov/nl	tococci, Aerococcus spp., and coccur within a timeframe that	Micrococcus spp.] is cultured from does not exceed a gap of 1





Central Line-Associated Bloodstream Infection (CLABSI): Mucosal Barrier Injury Laboratory Confirmed Bloodstream Infection (MBI-LCBI)

NHSN CLABSI protocol http://www.cdc.gov/nhsn/PDFs/pscManual/4PSC_CLABScurrent.pdf

Patient name _	MR #	Date of admission		
	l line insertion			Date of event
criteria may	be present during the first?	irst present together ≥ 3 calendar days of 2 calendar days of admission as long as the eria occur within a timeframe that does no	ey are also present on or after	calendar day 3 of admission
with day of		cal catheter (UC) in place for > 2 calendary 1(or, if the patient is admitted or transfer Day 3 of line	——————————————————————————————————————	<u> </u>
AND	•	•		
	C is in place on the day of event in place on day of event	rent or day before □ Line in place day before event		
Patient must m	neet at least one of the follow	ving criteria for mucosal barrier injury l	aboratory-confirmed bloods	stream infection (MBI-LCBI):
Criterion 1 (a	ny age)			
<u>organisms i</u> spp., <i>Veillo</i> i	• •	LCBI with at least one blood culture grow andida spp., Clostridium spp., Enterococa aceae*		<u> </u>
AND				
	ts at least one of the followi	=		Ui 1
0		etic stem cell transplant recipient within the	ie past year with one of the fol	llowing documented during same
	hospitalization as positive b	rointestinal graft versus host disease (GI C	SVHD)	
	$\circ \geq 1$ liter diarrhea in	a 24-hour period (or \geq 20 mL/kg in a 24-hore the date the positive blood culture was	nour period for patients < 18 y	ears of age) with onset on or within
0		t least 2 separate days with values of abso		
	$(WBC) < 500 \text{ cells/mm}^3 \text{ on}$	or within 3 calendar days before the date	the positive blood culture was	s collected (Day 1).

* Partial list of MBI-LCBI Criterion 1 eligible Enterobacteriaceae genera: Citrobacter, Enterobacter, Escherichia, Klebsiella, Proteus, Providencia,



Salmonella, Serratia, Shigella, Yersina



Central Line-Associated Bloodstream Infection (CLABSI): Mucosal Barrier Injury Laboratory Confirmed Bloodstream Infection (MBI-LCBI)

NHSN CLABSI protocol http://www.cdc.gov/nhsn/PDFs/pscManual/4PSC_CLABScurrent.pdf

Patient name	MR #	Date of admission		
Date of centra	l line insertion	Date of central line removal	Date of review	Date of event
present dur	ing the first 2 calendar of	are first present together ≥ 3 calendar days of a lays of admission as long as they are also present criteria occur within a timeframe that does not	nt on or after calendar day 3 c	of admission
device plac		nbilical catheter (UC) in place for > 2 calendar of the patient is admitted or transferred with a center of Day 3 of line		
AND	·	•		
	C is in place on the day			
□ Line	in place on day of even	t Line in place day before event		
Criterion 2 (a Patient of a AND		for LCBI when the blood cultures are growing	only viridans group streptoco	occi with no other organisms isolated
	ets at least one of the fol	owing:		
0		topoietic stem cell transplant recipient within th	e past year with one of the fo	llowing documented during same
		V gastrointestinal graft versus host disease (GI	GVHD)	
		nea in a 24-hour period (or \geq 20 mL/kg in a 24-late date the first positive blood culture was collected to the first positive blood culture was		years of age) with onset on or within 7 calendar
0		d as at least 2 separate days with values of abson 3 calendar days before the date the positive bl		
Criterion 3 (1	neonates or infants only	7)		
$□$ Patient ≤ 1 AND	year of age meets criter	on 3 for LCBI when the blood cultures are grow	ving only viridans group strep	otococci with no other organisms isolated
□ Patient mee	ets at least one of the fol	owing		
0	Is an allogeneic hema hospitalization as pos	topoietic stem cell transplant recipient within that it is blood culture:	e past year with one of the fo	llowing documented during same
		V gastrointestinal graft versus host disease (GI		
	$\circ \geq 20 \text{ mL/kg o}$ collected	f diarrhea in a 24-hour period with onset on or	within 7 calendar days before	the date the first positive blood culture was
0	3	d as at least 2 separate days with values of abso	_	





CLABSI Notes

- 1. In LCBI criterion 1, the phrase "one or more blood cultures" means that at least one bottle from a blood draw is reported by the laboratory as having grown at least one organism (i.e., is a positive blood culture).
- 2. In LCBI criterion 1, the term "recognized pathogen" does not include organisms considered common commensals (see criteria 2 and 3 for a list of common commensals). A few of the recognized pathogens *are S. aureus, Enterococcus* spp., *E. coli, Pseudomonas* spp., *Klebsiella* spp., *Candida* spp., etc.
- 3. In LCBI criteria 2 and 3, the phrase "two or more blood cultures drawn on separate occasions" means 1) that blood from at least two blood draws were collected within two calendar days of each other (e.g., blood draws on Monday and Tuesday would be acceptable for blood cultures drawn on separate occasions, but blood draws on Monday and Wednesday would be too far apart in time to meet this criterion), and 2) that at least one bottle from each blood draw is reported by the laboratory as having grown the same common commensal (i.e., is a positive blood culture). (See note 4 for determining sameness of organisms).
 - a. For example, an adult patient has blood drawn at 8 a.m. and again at 8:15 a.m. of the same day. Blood from each blood draw is inoculated into two bottles and incubated (four bottles total). If one bottle from each blood draw set is positive for coagulase-negative staphylococci, this part of the criterion is met.
 - b. For example, a neonate has blood drawn for culture on Tuesday and again on Thursday and both grow the same common commensal. Because the time between these blood cultures exceeds the two-day period for blood draws stipulated in LCBI and MBI-LCBI criteria 2 and 3, this part of the criterion is not met.
 - c. "Separate occasions" also means blood draws collected from separate sites or separate accesses of the same site, such as two draws from a single lumen catheter or draws from separate lumens of a catheter. In the latter case, the draws may be just minutes apart (i.e., just the time it takes to disinfect and draw the specimen from each lumen). For example, a patient with a triple lumen central line has blood drawn from each lumen within 15 minutes of each other. Each of them is considered a separate blood draw.
 - d. A blood culture may consist of a single bottle for a pediatric blood draw due to volume constraints. Therefore, to meet this part of the criterion, each bottle from two or more draws would have to be culture-positive for the same commensal.
- 4. If the pathogen or common commensal is identified to the species level from one blood culture, and a companion blood culture is identified with only a descriptive name (e.g., to the genus level), then it is assumed that the organisms are the same. The organism identified to the species level should be reported as the infecting organism along with its antibiogram if available.
- 5. Only genus and species identification should be utilized to determine the sameness of organisms (i.e., matching organisms). No additional comparative methods should be used (e.g., morphology or antibiograms) because laboratory testing capabilities and protocols may vary between facilities. This will reduce reporting variability, solely due to laboratory practice, between facilities reporting LCBIs meeting criterion 2. Report the organism to the genus/species level only once, and if antibiogram data are available, report the results from the most resistant panel.
- 6. LCBI criteria 1 and 2 and MCI-LCBI criteria 1 and 2 may be used for patients of any age, including these patients ≤1 year of age.
- 7. Specimen Collection Considerations: Although blood cultures drawn through central lines can have a higher rate of contamination than blood cultures collected through peripheral venipuncture, all positive blood cultures, regardless of the sites from which they were collected, must be included when conducting in-plan CLABSI surveillance.
- 8. "No other organisms isolated" means there is not isolation in a blood culture of another recognized pathogen (e.g., *S. aureus*) or common commensal (e.g., coagulase-negative staphylococci) other than listed in MBI-LCBI criterion 1, 2 or 3 that would otherwise meet LCBI criteria. If this occurs, the infection should not be classified as MBI-LCBI.
- 9. Grade III/IV GI GVHD is defined as follows:
 - a. In adults: ≥1 L diarrhea/day or ileus with abdominal pain
 - b. In pediatric patients: ≥20 cc/kg/day of diarrhea





CLABSI Reporting instructions:

- 1. Report organisms cultured from blood as BSI–LCBI when no other site of infection is evident (see Appendix 1. Secondary Bloodstream Infection (BSI) Guide).
- 2. Catheter tip cultures are not used to determine whether a patient has a primary BSI.
- 3. When there is a positive blood culture and clinical signs or symptoms of localized infection at a vascular access site, but no other infection can be found, the infection is considered a primary BSI.
- 4. Purulent phlebitis confirmed with a positive semi-quantitative culture of a catheter tip, but with either negative or no blood culture is considered a CVS-VASC, not a BSI or an SST-SKIN or ST infection.
- 5. Occasionally a patient with both peripheral and central IV lines develops a primary bloodstream infection (LCBI) that can clearly be attributed to the peripheral line (e.g., pus at the insertion site and matching pathogen from pus and blood). In this situation, enter "Central Line = No" in the NHSN application. You should, however, include the patient's central line days in the summary denominator count.
- 6. If your state or facility requires that you report healthcare-associated BSIs that are not central line-associated, enter "Central Line = No" in the NHSN application when reporting these BSIs. You should, however, include all of the patient's central line days in the summary denominator count.



